

# Pergoquin, 1 mg tablets for horses

Authorised

- Pergolide mesilate

## Product identification

**Medicine name:**

Pergoquin, 1 mg tablets for horses  
PERGOQUIN δισκία 1mg/tab

**Active substance:**

Pergolide mesilate

**Target species:**

Horse

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Pergolide mesilate  
1.31 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**

Tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN04BC02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Greece

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**Package description:**

Cardboard box of 200 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.

Cardboard box of 160 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.

Cardboard box of 150 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.

Cardboard box of 100 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.

Cardboard box of 50 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.

Cardboard box of 60 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Vetviva Richter GmbH

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**Marketing authorisation date:**

3/02/2020

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

11074/04-02-2020 K-0238701

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**Date of authorisation status change:**

3/02/2020

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0295/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Czechia Denmark Estonia Finland  
Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Norway  
Poland Portugal Romania Slovakia Slovenia Spain Sweden  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to

[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents